



Quality System Manual

MIPM - Brevier

International
Distributor
Questionnaire (IDQ)

II. SALES / MARKETING INFORMATION

Annual Revenue in € (last two years):

2018: GBP £2.3M ± €2.8M 2019: GBP £2.5M ± €3M

Distribution Network:

Total Company Employees:

21

Sales People:

8

Direct Sales force:

2

Service Engineer:

5

Administrative staff:

5

Areas Covered by Distribution Network:

UK AND IRELAND

WE ALSO EXPORT SOME PRODUCTS

Product Portfolio Currently Carried by Distribution Network:

Products:

OXYGEN SENSORS / MONITORS

FLOWSENSORS

PULSE OXIMETERS

AIR/OXYGEN BOTTLES

TEMPERATURE PROBES

Manufacturer:

TELEDYNE / INVITEC / ITGAMBSET / MAXTEC

BLUPOINT MEDICAL

BLUPOINT MEDICAL

MAXTEC

BLUPOINT MEDICAL

In which medical disciplines or hospital areas/departments are your representatives active?

Anaesthesia

Biomedical

Radiology

ICU

Pediatrics

NICU



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Which products are you interested in distributing for MIPM?

MRI Patient Monitoring: *Tesla*^{M3}
 MRI Patient Monitoring: *Tesla*^{DUO}
 TOF 3D.....

Are you able and willing to buy demonstration units?

Yes No

Are you able to perform repair and maintenance service for MIPM products?

Yes No

Are you able and willing to send personnel for training to Germany?

Yes No

Which medical congresses and trade shows are important in your territory?

WE ARE NOT BOUND TO EXHIBIT AT ANY CONFERENCES IN THE UK, AS WE HAVE FOUND IT MORE BENEFICIAL TO FOCUS OUR EFFORTS AND RESOURCES INTO DIRECT MARKETING/SELLING TO THE INDIVIDUAL HOSPITAL DEPARTMENTS.

Will you participate in it?

Yes No *NO PLANS, BUT WE EVALUATE THIS ON A REGULAR BASIS.*

If so, are you able and willing to show MIPM products at these shows?

Yes No *IF APPLICABLE*

How many hospitals are in your sales area?

< 50 < 100 < 200 > 200

How many MRI scanners are in your sales area (incl. private imaging centers)?

< 50 < 100 < 200 > 200

Which competitors are active in your sales area?

THERE IS CURRENTLY A GAP IN THE MARKET FOR A NERVE STIMULATOR, SINCE THE VIAMED MICROSIM WAS DISCONTINUED.



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III. REGULATORY AFFAIRS

Is a regulatory approval required in your country or distribution region?

Yes No

If so, is CE-Certification or FDA approval sufficient?

Yes No

If not, are you able and willing to take responsibility for regulatory affairs according to the regulatory requirements in your country?

Yes No

If so, please provide your Importer Establishment Registration Number:

Questionnaire filled by

Name & Title: *RYAN SWAIN - INTERNATIONAL SALES MANAGER*

Date: *17/2/2020*



→ Please return filled out form to: international@mipm.com